

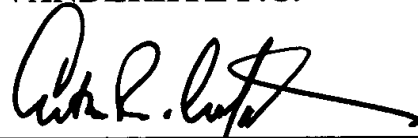
REMARKS

Attached hereto is a marked-up version of the changes made to the specification and claims by the current amendment. The attached page(s) is captioned "**Version With Markings To Show Changes Made.**"

Respectfully submitted,

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VERSION WITH MARKINGS TO SHOW CHANGES MADE

IN THE SPECIFICATION

Page 1, before the first line, insert as a separate paragraph:

This application is the US national phase of international application
PCT/EP00/09708 filed 4 October 2000, which designated the US.

IN THE CLAIMS

11. The composition of ~~either of claims 5 or 9~~ wherein the agonist is used to treat endometriosis.

12. The composition of ~~either of claims 5 or 9~~ wherein the agonist is used to treat infertility or to improve fertility.

20. The method of ~~either of claims 15 or 19~~, wherein the composition is administrated every 12 to 96 hours.

21. The method of ~~either of claims 15 or 19~~, wherein composition is administrated twice weekly.

22. Te method of ~~either of claims 15 or 19~~, wherein the composition further comprises a tablet.

25. Use accordingly to claim 23-~~or~~ 24 wherein the medicament is formulated to deliver from less than 1 mg to 8 mg of the β -adrenergic agonist per dose.

26. Use according to ~~any one of claim 23 to 25~~ wherein the β -adrenergic agonist is terbutaline.

27. Use according to ~~any one of claims 23 to 26~~ wherein the medicament is formulated to deliver from less than 1 mg to 8 mg of the β -adrenergic agonist per dose.

28. Use according to ~~any one of claims 23 to 27~~ wherein the bioadhesive carrier comprises a cross-linked water-insoluble but water-swellaable polycarboxylic acid polymer.

32. Use according to claim 30-~~or~~ 31 wherein the medicament is for administration every 12 to 96 hours.

33. Use according to claim 30-~~or~~ 31 wherein the medicament is for administration twice weekly.

34. Use according to ~~any one of claims 23 to 33~~ wherein the medicament avoids detrimental blood levels of the β -adrenergic agonist.

